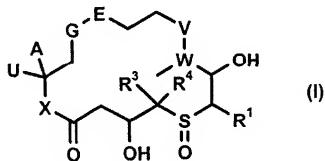


# AMENDMENTS TO THE CLAIMS

Please amend the claims as follows.

## 1. (Previously Presented) Compounds of Formula (I)

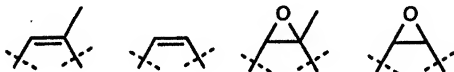


wherein

A is a group of the formula  $-\text{C}(\text{CH}_3)=\text{CHR}^5$  or  $-\text{CH}=\text{CHR}^5$ , wherein  $\text{R}^5$  is a heteroaryl- or a heteroarylalkyl group,

U is hydrogen, halogen,  $\text{C}_1$ - $\text{C}_4$  alkyl,  $\text{C}_3$ - $\text{C}_4$ -cycloalkyl,  $\text{C}_1$ - $\text{C}_4$  heteroalkyl-, trifluoromethyl or  $\text{COOH}$ ,

G-E is selected from the following groups,



or is part of an optionally substituted phenyl ring,

$R^1$  is a  $C_1$ - $C_4$ -alkyl-, a  $C_2$ - $C_4$ -alkenyl-, a  $C_2$ - $C_4$ -alkynyl- or a  $C_3$ - $C_4$ -cycloalkyl-group,

V-W is a group of formula  $CH_2CH$  or  $CH=C$ ,

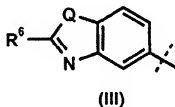
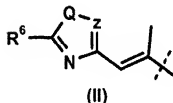
X is oxygen or a group of the formula  $NR^2$ , wherein  $R^2$  is hydrogen,  $C_1$ - $C_4$  alkyl,  $C_2$ - $C_4$  alkenyl, or  $C_1$ - $C_4$  heteroalkyl,  
and

$R^3$  and  $R^4$  independently from each other represent hydrogen,  $C_1$ - $C_4$ -alkyl or together are part of a cycloalkyl group with 3 or 4 ring atoms,

or a pharmacologically acceptable salt, solvate, hydrate or formulation thereof.

2. (Canceled)

3. (Original) Compounds according to claim 1, wherein A is a group of formula (II) or (III)



wherein Q is sulphur, oxygen or  $NR^7$ , wherein  $R^7$  is hydrogen,  $C_1$ - $C_4$  alkyl or  $C_1$ - $C_4$  heteroalkyl, Z is Nitrogen or CH and  $R^6$  is  $OR^8$ ,  $NHR^8$ ,  $C_1$ - $C_4$  alkyl,  $C_2$ - $C_4$  alkenyl,  $C_2$ - $C_4$  alkynyl or  $C_1$ - $C_6$  heteroalkyl, wherein  $R^8$  is hydrogen,  $C_1$ - $C_4$  alkyl or  $C_1$ - $C_4$  heteroalkyl.

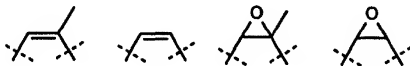
4. (Previously Presented) Compounds according to claim 1, wherein X is oxygen or NH.

5. (Previously Presented) Compounds according to claim 1, wherein  $R^1$  is methyl or ethyl.

6. **(Previously Presented)** Compounds according to claim 1, wherein  $R^3$  and  $R^4$  are methyl groups.

7. **(Previously Presented)** Compounds according to claim 1, wherein U is hydrogen, fluorine, methyl, trifluoromethyl or COOH.

8. **(Previously Presented)** Compounds according to claim 1, wherein G-E is selected from the following groups:



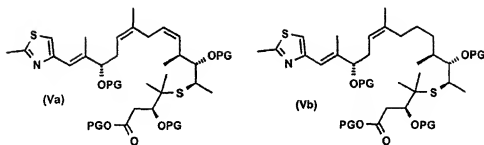
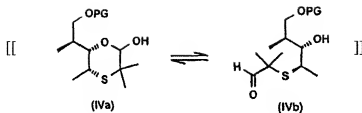
9. **(Previously Presented)** Compounds according to claim 1, wherein V-W is  $CH_2CH$ .

10. **(Previously Presented)** Pharmaceutical compositions containing a compound, a pharmacologically acceptable salt, a solvate or a hydrate according to claim 1 or a prodrug of the compound, the salt, the solvate and/or the hydrate and optionally one of more carriers and/or one or more adjuvants and/or one or more diluents.

11. **(Previously Presented)** Method of treating a disease selected from the group consisting of breast, ovarian, lung and prostate cancer through administering a pharmaceutically effective amount of a compound or a pharmaceutical composition according to claim 1.

12. **(Canceled)**

13. **(Currently Amended)** Compounds of formula ~~(IVa)~~, ~~(IVb)~~, (Va) and (Vb),



wherein the groups PG independently from each other represent hydrogen or protecting groups.

14. **(Currently Amended)** A process for preparing a compound of formula (I), comprising reacting Use of a compound according to claim 13 for the synthesis of a compound of formula (I) by

- a) removing any protecting groups on the acid and allyl alcohol;
- b) lactonizing the hydroxy acid;
- c) removing any protecting groups on the remaining alcohols;
- d) reducing the disubstituted double bond, if present; and
- e) oxidizing the sulfur atom at the 5-position to a sulfoxide.